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International Buréau INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT) (51) International Patent Classification 7: WO 00/23057 (11) International Publication Number: A2 A61K 31/00 (43) International Publication Date: 27 April 2000 (27.04.00) PCT/EP99/07804 (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, (21) International Application Number: BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, (22) International Filing Date: 12 October 1999 (12.10.99) KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, (30) Priority Data: UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, 98203454.8 16 October 1998 (16.10.98) EP MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, (71) Applicant (for all designated States except US): JANSSEN NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, PHARMACEUTICA N.V. [BE/BE]; Patent Department, GN, GW, ML, MR, NE, SN, TD, TG). Turnhoutseweg 30, B-2340 Beerse (BE). (72) Inventors; and (75) Inventors/Applicants (for US only): DE NIJS, Paul, Leonce, Published Without international search report and to be republished Ima [BE/BE]; (BE). PARYS, Wim, Louis, Julien [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 upon receipt of that report. Beerse (BE). (74) Agent: QUAGHEBEUR, Luc; Janssen Pharmaceutica N.V., Patent Department - ext. 3547, Turnhoutseweg 30, B-2340 Beerse (BE). (54) Title: THERAPY FOR IMPROVING COGNITION (57) Abstract The present invention is concerned with pharmaceutical compositions comprising a carrier and as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), each in an amount producing a therapeutically beneficial effect in patients suffering from psychosis, or Alzheimer's disease or related dementias. Said therapeutically beneficial effect can be a synergistic effect on the cognitive functioning of patients suffering from Alzheimer's disease or related dementias or the prevention of the further deterioration of cognition in said patients, or the reduction of adverse effects associated with the one of the active ingredients by the other of the active ingredients.

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THERAPY FOR IMPROVING COGNITION

The present invention is concerned with pharmaceutical compositions comprising a carrier and as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), each in an amount producing a therapeutically beneficial effect in patients suffering from psychosis, or Alzheimer's disease or related dementias. Said therapeutically beneficial effect can be a synergistic effect on the cognitive functioning of patients suffering from Alzheimer's disease or related dementias, or the prevention of the further deterioration of cognition in said patients, or the reduction of adverse effects associated with the one of the active ingredients by the other of the active ingredients.

Of particular interest is the use of an atypical antipsychotic agent (I) for the preparation of a medicament for reducing adverse effects associated with acetylcholinesterase inhibitors (II) in patients suffering from Alzheimer's disease or related dementias, such as nausea, vomiting, sweating, restlessness and insomnia. Especially interesting is the use of an atypical antipsychotic agent (I) for the preparation of a medicament for improving sleep in patients suffering from Alzheimer's disease or related dementias while being treated with acetylcholinesterase inhibitors (II).

The present invention is concerned with a pharmaceutical composition comprising a carrier and as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), each in an amount producing a therapeutically beneficial effect in patients suffering from psychosis, or Alzheimer's disease or related dementias. Said therapeutically beneficial effect can be a synergistic effect on the cognitive functioning of patients suffering from Alzheimer's disease or related dementias, or the prevention of the further deterioration of cognition in said patients, or the reduction of adverse effects associated with the one of the active ingredients by the other of the active ingredients.

The atypical antipsychotic (I) is selected from risperidone, 9-hydroxyrisperidone or a C₁₀₋₂₀ alkanoic acid ester thereof, olanzapine, quetiapine, iloperidone or ziprasidone, and the acetylcholinesterase inhibitor (II) is selected from galantamine, rivastigmine or donepezil, or therapeutically active acid addition salt form of any of the foregoing. Said salts comprise salt forms which the active ingredients (I) and (II) are able to form with appropriate acids, such as, for example, inorganic acids such as hydrohalic acids, e.g. hydrochloric or hydrobromic acid; sulfuric; nitric; phosphoric and the like acids; or

organic acids such as, for example, acetic, propanoic, hydroxyacetic, lactic, pyruvic, oxalic, malonic, succinic, maleic, fumaric, malic, tartaric, citric, methanesulfonic, ethanesulfonic, benzenesulfonic, p-toluenesulfonic, cyclamic, salicylic, p-aminosalicylic, pamoic and the like acids. For example, galantamine may conveniently be used as the (1:1) hydrobromide salt.

C10-20alkanoic acids are selected from the group consisting of decanoic (capric), undecanoic, dodecanoic (lauric), tridecanoic, tetradecanoic (myristic), pentadecanoic, hexadecanoic (palmitic), heptadecanoic, octadecanoic (stearic), nonadecanoic and eicosanoic acid. Due to their limited aqueous solubility, it was generally believed that the esters had to be suspended into oils. The ester having a C15 (pentadecyl) chain and the active ingredient corresponding thereto being the 9-hydroxyrisperidone palmitate ester was found to be the superior ester from a pharmacokinetic, as well as from a tolerance point of view.

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Preferably, the amount of each of the active ingredients is equal to or less than that which is approved in monotherapy with said active ingredient.

Most preferred are compositions wherein the atypical antipsychotic (I) is risperidone and the acetylcholinesterase inhibitor (II) is galantamine, in particular as galantamine hydrobromide. In said compositions, the amount of risperidone is 0.5, 1, 2, 4, or 6 mg and that of galantamine (as base) is 8, 16, 24 or 32 mg per dosage form.

The present invention also relates to products containing as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), as combined preparations for simultaneous, separate or sequential use in the treatment of patients suffering from psychosis, Alzheimer's disease or related dementias.

The present invention also concerns the use of an acetylcholinesterase inhibitor (II) for the preparation of a medicament for enhancing the effect of an atypical antipsychotic agent (I) on cognition in patients suffering from psychosis.

Conversely, the present invention also concerns the use of an atypical antipsychotic
agent (I) for the preparation of a medicament for enhancing the effect of an
acetylcholinesterase inhibitor (II) on cognition in patients suffering from Alzheimer's
disease or related dementias.

Additionally, the present invention concerns the use of an atypical antipsychotic agent (I) for the preparation of a medicament for reducing adverse effects associated with acetylcholinesterase inhibitors (II) in patients suffering from Alzheimer's disease or related dementias. Said adverse effect can be nausea, vomiting, sweating, restlessness or insomnia. Especially interesting is the use of an atypical antipsychotic agent (I) for the preparation of a medicament for improving sleep in patients suffering from Alzheimer's disease or related dementias while being treated with acetylcholinesterase inhibitors (II).

- Finally, the present invention also concerns the use of an acetylcholinesterase inhibitor (II) for the preparation of a medicament for reducing adverse effects associated with atypical antipsychotic agents (I) in patients suffering from psychoses. Said the adverse effect can be extrapyramidal syndrome or tardive dyskinesia.
- In all the preceding uses the atypical antipsychotic (I) is preferably risperidone and the acetylcholinesterase inhibitor (II) is preferably galantamine, in particular the (1:1) hydrobromide.

Claims

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- 1. A pharmaceutical composition comprising a carrier and as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), each in an amount producing a therapeutically beneficial effect in patients suffering from psychosis, Alzheimer's disease or related dementias.
- 2. A composition according to claim 1 wherein said therapeutically beneficial effect is a synergistic effect on the cognitive functioning of patients suffering from Alzheimer's disease or related dementias, or the prevention of the further deterioration of cognition in said patients, or the reduction of adverse effects associated with the one of the active ingredients by the other of the active ingredients.
 - 3. A composition according to claim 1 wherein the atypical antipsychotic (I) is selected from risperidone, 9-hydroxyrisperidone or a C₁₀₋₂₀ alkanoic acid ester thereof, olanzapine, quetiapine, iloperidone or ziprasidone, and the acetylcholinesterase inhibitor (II) is selected from galantamine, rivastigmine or donepezil.
 - 4. A composition according to claim 3 wherein the amount of each of the active ingredients is equal to or less than that which is approved in monotherapy with said active ingredient.
 - 5. A composition according to claim 3 wherein the atypical antipsychotic (I) is risperidone and the acetylcholinesterase inhibitor (II) is galantamine.
- 6. A composition according to claim 5 wherein the amount of risperidone is 0.5, 1, 2, 4, or 6 mg and that of galantamine (as base) is 8, 16, 24 or 32 mg per dosage form.
 - 7. A product containing as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), as a combined preparation for simultaneous, separate or sequential use in the treatment of patients suffering from psychosis, Alzheimer's disease or related dementias.

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- 8. The use of an acetylcholinesterase inhibitor (II) for the preparation of a medicament for enhancing the effect of an atypical antipsychotic agent (I) on cognition in patients suffering from psychosis.
- 5 9. The use of an atypical antipsychotic agent (I) for the preparation of a medicament for enhancing the effect of an acetylcholinesterase inhibitor (II) on cognition in patients suffering from Alzheimer's disease or related dementias.
- 10. The use of an atypical antipsychotic agent (I) for the preparation of a medicament for reducing adverse effects associated with acetylcholinesterase inhibitors (II) in patients suffering from Alzheimer's disease or related dementias.
 - 11. Use according to claim 10 wherein the adverse effect is nausea, vomiting, sweating, restlessness or insomnia.
 - 12. The use of an acetylcholinesterase inhibitor (II) for the preparation of a medicament for reducing adverse effects associated with atypical antipsychotic agents (I) in patients suffering from psychoses.
- 20 13. Use according to claim 12 wherein the adverse effect is extrapyramidal syndrome or tardive dyskinesia.
 - 14. Use according to any one of claims 8 to 13 wherein the atypical antipsychotic (I) is risperidone and the acetylcholinesterase inhibitor (II) is galantamine.

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(72) Inventors; and

- (75) Inventors/Applicants (for US only): DE NIJS, Paul, Leonce, Irma [BE/BE]; (BE). PARYS, Wim, Louis, Julien [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE).
- (74) Agent: QUAGHEBEUR, Luc; Janssen Pharmaceutica N.V. Patent Department - ext. 3547, Turnhoutseweg 30, B-2340 Beerse (BE).

KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS. MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

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With international search report.

(88) Date of publication of the international search report: 27 July 2000 (27.07.00)

(54) Title: ATYPICAL ANTIPHSYCHOTIC IN COMBINATION WITH ACETYLCHOLINESTERASE INHIBITOR FOR IMPROVING COGNITION

(57) Abstract

The present invention is concerned with pharmaceutical compositions comprising a carrier and as first active ingredient an atypical antipsychotic agent (I) and as second active ingredient an acetylcholinesterase inhibitor (II), each in an amount producing a therapeutically beneficial effect in patients suffering from psychosis, or Alzheimer's disease or related dementias. Said therapeutically beneficial effect can be a synergistic effect on the cognitive functioning of patients suffering from Alzheimer's disease or related dementias or the prevention of the further deterioration of cognition in said patients, or the reduction of adverse effects associated with the one of the active ingredients by the other of the active ingredients.

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inter onal Application No PCT/EP 99/07804

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 A61K31/55

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) $IPC\ 7 \ A61K$

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

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•	EP 0 879 596 A (SNORASSON ERNIR) 25 November 1998 (1998-11-25) abstract; claims 37,38 page 3	1-14
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E	WO 99 52519 A (GEN HOSPITAL CORP) 21 October 1999 (1999-10-21) claims 34,37,38 -/	1-16

X Further documents are listed in the continuation of box C.	Patent family members are listed in annex.
Special categories of cited documents: A document defining the general state of the art which is not considered to be of particular relevance. E earlier document but published on or after the international filing date. L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified). O document referring to an oral disclosure, use, exhibition or other means. P document published prior to the international filing date but later than the priority date claimed.	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention. "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone. "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search 4 April 2000	Date of mailing of the international search report $26/04/2000$
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Gonzalez Ramon, N

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Intel onal Application No PCT/EP 99/07804

C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	FC1/EF 99/0/804
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	WO 99 07378 A (SANBERG PAUL RONALD ;SHYTLE ROLAND DOUGLAS (US); SILVER ARCHIE AAR) 18 February 1999 (1999-02-18) abstract page 8 -page 9; claim 11	1-16
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T	MAZUREK A. A.: "Treatment of Alzheimer's disease" N ENG J. MED, vol. 342, no. 11, 16 March 2000 (2000-03-16), pages 821-822, XP000901026 page 821, column 2, paragraph 1	1-16
Х,Р	YAQUB B.A.: "New horizons in management of Alzheimer 's disease." SAUDI MEDICAL JOURNAL, (1999) 20/9 (671-677)., XP000892838 page 672 -page 673; table 1	1-16

Intel onal Application No PCT/EP 99/07804

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Υ,Ρ	MELTZER, H. Y. (1) ET AL: "Atypical antipsychotic drugs (APD) but not typical APD increased extracellular acetylcholine (ACh-ext) levels in rat medial prefrontal cortex (mPFC) in the absence AChEsterase (AChE) inhibition." SOCIETY FOR NEUROSCIENCE ABSTRACTS, (1999) VOL. 25, NO. 1-2, PP. 452. MEETING INFO.: 29TH ANNUAL MEETING OF THE SOCIETY FOR NEUROSCIENCE, PART 1 MIAMI BEACH, FLORIDA, USA OCTOBER 23-28, 1999 THE SOCIETY FOR NEUROSCIENCE., XPO00892699 abstract	1-16
X	MAGNUSON T M ET AL: "Extrapyramidal side effects in a patient treated with risperidone plus donepezil 'letter!." AMERICAN JOURNAL OF PSYCHIATRY, (1998 OCT) 155 (10) 1458-9. , XP000892634 the whole document	1-16
X	SIMONSON W.: "Promising agents for treating Alzheimer's disease." AMERICAN JOURNAL OF HEALTH-SYSTEM PHARMACY, (1 NOV 1998) 55/21 SUPPL. (S11-S16)., XP002134583 page S14, column 2, paragraph 3	1-16
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In...mational application No.

PCT/EP 99/07804

Box i Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
Claims Nos.: 1,2,7-13 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically: SEE FURTHER INFORMATION SHEET PCT/ISA/210
Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II Observations where unity of Invention is lacking (Continuation of Item 2 of first sheet)
This International Searching Authority found multiple inventions in this international application, as follows:
As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest The additional search fees were accompanied by the applicant's protest. No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet (1)) (July 1998)

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1,2,7-13

Present claims 1,2,7-13 relate to compounds defined by reference to the following parameters: "atypical antipsychotic agent" and "acetylcholinesterase inhibitor".

The use of these parameters in the present context is considered to lead to a lack of clarity within the meaning of Article 6 PCT. It is impossible to compare the parameters the applicant has chosen to employ with what is set out in the prior art. The lack of clarity is such as to render a meaningful complete search impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds specifically disclosed in the examples and the claims with due regard to the general idea underlying the present application

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

information on patent family members

Intel onal Application No PCT/EP 99/07804

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